

**DRAFT**  
**ENHANCED BIOREMEDIATION FIELD TEST REPORT**  
**OPERABLE UNIT 2**  
**SITE ST012 – LIQUID FUELS STORAGE AREA**  
**FORMER WILLIAMS AIR FORCE BASE**  
**MESA, ARIZONA**

**Prepared for:**  
**Air Force Civil Engineer Center**  
**AFCEC/CIBW**  
**2261 Hughes Avenue, Suite 155**  
**Lackland AFB, Texas 78236-9851**

**Prepared by:**  
**Amec Foster Wheeler Environment & Infrastructure, Inc.**  
**4600 East Washington Street, Suite 600**  
**Phoenix, Arizona 85034**

**30 November 2015**

**Contract Number: FA8903-09-D-8572 - 0002**  
**Project No. 9101110001**  
**CDRL No. A001**

**SIGNATURE PAGE**

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%	percent
AFB	Air Force Base
AFCEC	Air Force Civil Engineer Center
Amec Foster Wheeler	Amec Foster Wheeler Environment & Infrastructure, Inc.
CZ	Cobble Zone
DNF	denitrifying bacteria
EBR	enhanced bioremediation
frac tank	fractionation tank
ft msl	feet above mean sea level
GEO	geobacter
gpm	gallons per minute
JP-4	jet propulsion fuel grade 4
LNAPL	light non-aqueous phased liquid
LSZ	Lower Saturated Zone
mg/l	milligram per liter
MGN	methanogenic bacteria
Microbial Insights	Microbial Insights, Inc.
OU-2	Operable Unit 2
qPCR	quantitative polymerase chain reaction
PLFA	phospholipid fatty acid
RD/RAWP	Remedial Design/Remedial Action Work Plan
RODA-2	Record of Decision Amendment 2
SEE	Steam Enhanced Extraction
SRB	sulfate reducing bacteria
ST012	Site ST012 Liquid Fuel Storage Area
TEA	terminal electron acceptor
TestAmerica	TestAmerica Laboratories, Inc.
TPH	total petroleum hydrocarbons
TTZs	Thermal Treatment Zones
UWBZ	Upper Water Bearing Zone
µg/l	microgram per liter
VOC	volatile organic compound

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## **1.0 INTRODUCTION**

Amec Foster Wheeler Environment & Infrastructure, Inc. (Amec Foster Wheeler) prepared this Enhanced Bioremediation (EBR) Field Test Report for the Air Force Civil Engineer Center (AFCEC) to provide details related to EBR field activities conducted for Operable Unit 2 (OU-2) at the former Williams Air Force Base (AFB) in Mesa, Arizona (Figure 1-1). OU-2 is composed of the Installation Restoration Program Site ST012, the former Liquid Fuels Storage Area (ST012) (Figure 1-2), where historic releases of jet propulsion fuel grade 4 (JP-4) and/or aviation gasoline have resulted in contaminated soil and groundwater.

An OU-2 Record of Decision Amendment 2 (RODA 2) (AFCEC, 2013) revised the groundwater remedy to implement steam enhanced extraction (SEE) followed by EBR. The purpose of the EBR Field Test was to obtain data to support EBR design.

### **1.1 Purpose and Organization of Report**

The purpose of this EBR Field Test Completion Report is to document the field activities and associated results related to the EBR Field Test Plan (AMEC, 2014b), including interpretation of results.

This report consists of five sections, plus appendices. Section 1.0 presents an introduction and background information. Section 2.0 describes the field activities including deviations from the Final EBR Field Test Plan (AMEC, 2014b) and laboratory results. Section 3.0 discusses results and estimates parameters for use in EBR design. Appendix A presents low-flow sample collection records. Appendix B includes analytical laboratory results. Data validation narratives are included in Appendix C and microbial results are in Appendix D. Appendix E presents calculations for the estimate of microbial degradation kinetics.

### **1.2 Background**

As part of the ST012 Remedial Design and Remedial Action Work Plan (RD/RAWP) (AMEC, 2014a) for implementing the OU-2 RODA 2, the selected remedial action includes an initial period of SEE for mass removal of dissolved contaminants and light non-aqueous phased liquid (LNAPL) within established thermal treatment zones (TTZs), followed by EBR to address LNAPL outside of the TTZs as well as dissolved phase contaminants within and outside the TTZs. Amec Foster Wheeler conducted this EBR Field Test in support of the design for the enhanced bioremediation portion of the approved groundwater remedy for the site. The EBR and SEE modeling report prepared as part of the ST012 RD/RAWP described EBR modeling results using both aerobic and anaerobic processes for remediation. Modeling results for both processes indicated that the remediation goals could be met; however, there were significant assumptions made for the anaerobic approach. Therefore, this EBR Field Test included two single well tracer tests, herein also referred to as push-pull tests, using sulfate as a terminal electron acceptor (TEA) to evaluate appropriate delivery and dosing for EBR at ST012 under the anaerobic scenario. Due to the prevalence of existing aerobic EBR data that can adequately support a future design, this field test only investigated anaerobic EBR to provide data to support evaluation of strategies for

aerobic and/or anaerobic EBR and for design of anaerobic methods if this approach is implemented.

Monitoring wells ST012-W11 and ST012-W30 were selected for the single-well tracer tests because they are both located outside of the SEE TTZs and both have historical evidence of LNAPL contamination. Both wells are screened in the Lower Saturation Zone (LSZ) and are located on the United States Army Reserve Property to the west of the SEE TTZs (see Figure 1-3). Figure 1-3 also shows the extent of residual LNAPL within each hydrostratigraphic zone as estimated in the RD/RAWP (AMEC, 2014a).

### **1.3 Field Test Procedure**

The EBR field test was conducted as two separate push-pull tests at well locations ST012-W11 and ST012-W30. The field test was conducted in general conformance with the EBR Field Test Plan (AMEC, 2014b) which included the following steps:

1. Baseline Sampling and Well Preparation
2. TEA and tracer (solution) mixing and injection
3. Monitoring during shut-in period
4. Groundwater/solution extraction and sampling during extraction
5. Post-extraction sampling
6. Review and interpretation of laboratory data

Each of the above steps is discussed in detail in Sections 2 through 4.

## 2.0 EBR FIELD TEST IMPLEMENTATION

The EBR Field Test was conducted in general accordance with the Final EBR Field Test Plan (AMEC, 2014b). This section describes implementation activities.

### 2.1 Site Preparation

Coordination with the U.S. Army Reserve Center occurred to gain access to the site for the test and agree on locations for staging of equipment and tanks. Equipment required for implementation was temporarily located within the U.S. Army Reserve Center during the push and pull phases of the EBR test except for the fractionation tank (frac tank) which was used to store extracted groundwater during the pull phase and was located on Avoca Street (which was closed for SEE construction activities).

### 2.2 Baseline Sampling and Well Preparation

Prior to initiating the field test, baseline samples were collected. On 16 July 2014, low-flow procedures were followed to collect samples for baseline analysis from monitoring wells ST012-W11 and ST012-W30. Once field readings of temperature, pH, specific conductance, dissolved oxygen, and oxidation-reduction potential stabilized (see Appendix A), samples were collected and submitted to Test America Laboratories (Test America) and Microbial Insights, Inc. (Microbial Insights) for the following analyses:

- Chemical Analysis (Test America):
  - sulfate
  - bromide
  - cation/anion balance
    - Inductively Coupled Plasma Metals
    - Ion Chromatography
  - Total Petroleum Hydrocarbons (TPH)
  - Volatile Organic Compounds (VOCs)
- Microbial Population Analysis (Microbial Insights):
  - quantitative polymerase chain reaction (qPCR) analysis to enumerate the following bacterial groups:
    - denitrifying bacteria (DNF, nirK and nirS functional genes)
    - geobacter (GEO)
    - sulfate reducing bacteria (SRB; APS)
    - methanogenic bacteria (MGN)
  - phospholipid fatty acid (PLFA) analysis to quantify total biomass and assess microbial population activity

On 16 July 2014, attempts were made to use an in-line bio-flo™ filter provided by Microbial Insights to collect samples for microbial population analysis; however, the filters clogged before the required amount of water was collected. Microbial Insights indicated prior to the field test that this is a common occurrence and provided glass bottles to fill with water instead of using the filters



if this problem arose. When the samples arrived at Microbial Insights, the laboratory indicated that there was not enough sample to run both sets of tests, so additional samples were collected from each well on 17 July 2014 to conduct the PLFA analysis.

Prior to beginning the field tests, an oil-water interface probe was used to check for LNAPL in the wells. A trace of LNAPL was detected in ST012-W11 but there was not enough to remove with a bailer. A pressure transducer and AquiStar® TempHion™ Smart Sensor (an in-situ bromide probe and data logger) were then placed inside each well to monitor changes in groundwater elevation and bromide concentrations, respectively, during the testing.

### **2.2.1 Analytical Data from Test America**

Tables 2-1 and 2-2 present analytical data from Test America for the full suite of analyses performed during baseline, post-shut-in, and post-extraction sampling at ST012-W11 and ST012-W30, respectively. Analytical reports are included in Appendix B and data validation narratives are provided in Appendix C. Sections 2.4 and 2.5 provide additional details on sample collection activities during the post-shut-in and post-extraction periods, respectively.

**Table 2-1 Analytical Data for ST012-W11**

Sample Location: Sample Type: Sample Date: Parameter		ST012 - W11			
		Baseline 7/16/2014	Post-Shut-In 9/2/2014	Post-Extraction 9/8/2014	
	Units				
JP-4 (Total Petroleum Hydrocarbons)					
Diesel Range Organics [C10-C28]	mg/l	0.45 B	0.56	0.095	U
Gasoline Range Organics [C6-C10]	µg/l	3000	2700 Q	73	Q
<i>Total Petroleum Hydrocarbons</i>	<i>mg/l</i>	3.5	3.3	0.17	
Volatile Organic Compounds					
1,2-Dichloroethane	µg/l	0.4 U	0.4 UQ	0.4	U
Benzene	µg/l	51	45 Q	0.37	J
Ethylbenzene	µg/l	310	230	2.4	
Methylene Chloride	µg/l	0.8 U	1.1 J	0.8	U
m-Xylene & p-Xylene	µg/l	0.8 U	74 D	2.2	
Naphthalene	µg/l	38 B	43	0.4	
o-Xylene	µg/l	0.4 U	4 U	0.4	U
Toluene	µg/l	0.38 J	1.2 Q	0.4	U
Trichloroethene	µg/l	0.21 J	0.2 UQ	0.2	U
Trichlorofluoromethane	µg/l	0.8 U	0.8 U	0.8	U
Xylenes, Total	µg/l	1.6 U	74 D	2.2	
<i>Total Volatile Organic Compounds</i>	<i>mg/l</i>	0.40	0.47	0.011	
Other					
Bromide	mg/l	1.6 J	2.6	2.5	
Sulfate	mg/l	5.4	6.4	280	
Calcium	mg/l	310	300	770	
Chloride	mg/l	800	780	1600	
Iron	mg/l	0.063 J	0.14	0.03	U
Manganese	mg/l	2.2 Q	2 Q	0.067	Q
Magnesium	mg/l	67	67	140	
Sodium	mg/l	79	76	92	
Potassium	mg/l	7.7	16	21	

**Notes:**

JP-4 – jet propulsion fuel grade 4

µg/l – micrograms per liter

mg/l – milligrams per liter

B = The analyte was detected above one-half the reporting limit in an associated blank

J = The analyte was positively identified; the quantitation is an estimation

Q = One or more quality control criteria failed

**Table 2-2 Analytical Data for ST012-W30**

Sample Location: Sample Type: Sample Date:		ST012 - W30					
		Baseline 7/16/2014		Post-Shut-In 9/2/2014		Post-Extraction 9/10/2014	
Parameter	Units						
JP-4 (Via Petroleum Hydrocarbons)							
Diesel Range Organics [C10-C28]	mg/l	0.47	B	1.2		11	
Gasoline Range Organics [C6-C10]	µg/l	3900		13000		14000	
Total Petroleum Hydrocarbons	mg/l	4.37		14.2		25	
Volatile Organic Compounds							
1,2-Dichloroethane	µg/l	4	U	4	U	2	U
Benzene	µg/l	1200		2700		1900	
Ethylbenzene	µg/l	290		920		740	
Methylene Chloride	µg/l	8	U	12	J	4	U
m-Xylene & p-Xylene	µg/l	8	U	570		950	
Naphthalene	µg/l	32	B	110		84	
o-Xylene	µg/l	4	U	130		85	
Toluene	µg/l	4	U	160		160	
Trichloroethene	µg/l	2	U	2	U	1	U
Trichlorofluoromethane	µg/l	8	U	8	U	4	U
Xylenes, Total	µg/l	16	U	700		1000	
Total Volatile Organic Compounds	mg/l	1.576		5.316		4.93	
Other							
Bromide	mg/l	1.3		3.5	J	2.3	
Sulfate	mg/l	11		18		13	
Calcium	mg/l	240		190		200	
Chloride	mg/l	600		580		510	
Iron	mg/l	0.65		1.2		2.7	
Manganese	mg/l	2.6	Q	3.3	Q	3.2	Q
Magnesium	mg/l	52		43		43	
Sodium	mg/l	67		62		60	
Potassium	mg/l	6.7		91		55	

**Notes:**

JP-4 – jet propulsion fuel grade 4  
 µg/l – micrograms per liter  
 mg/l – milligrams per liter

**Data Validation Flags:**

B – The analyte was detected above one-half the reporting limit in an associated blank  
 J – The analyte was positively identified; the quantitation is an estimation  
 Q – One or more quality control criteria failed

**2.2.2 Microbial Data from Microbial Insights**

Tables 2-3 and 2-4 present microbial data from Microbial Insights for the testing conducted during baseline, post-shut-in, and post-extraction sampling at ST012-W11 and ST012-W30, respectively. Microbial Analytical Reports are included in Appendix D.

**Table 2-3 Microbial Data for ST012-W11**

Sample Location:		ST012 - W11		
Parameter	Sample Type:	Baseline 7/16/2014 7/17/2014 <sup>1</sup>	Post-Shut-In 9/2/2014	Post-Extraction 9/8/2014
	Units			
qPCR				
Denitrifying Bacteria (nirK)	cells/ml	2.26E+06	4.52E+05	5.93E+05
Denitrifying Bacteria (nirS)	cells/ml	1.34E+05	1.93E+04	4.78E+04
Geobacter (GEO)	cells/ml	8.94E+01	1.53E+04	7.88E+02
Sulfate Reducing Bacteria (APS)	cells/ml	3.76E+04	6.49E+05	2.76E+06
Methanogens (MGN)	cells/ml	5.73E+05	7.00E+04	4.41E+03
PLFA				
Total Biomass	cells/ml	2.82E+04	4.39E+04	4.60E+04
Firmicutes	% of Total	34.68	20.44	0.52
Proteobacteria	% of Total	33.82	45.98	76.03
Anaerobic Metal Reducers	% of Total	1.68	1.91	0
SRB/Actinomycetes	% of Total	2.77	3.21	0.48
General	% of Total	27.06	27.09	22.42
Eukaryotes	% of Total	0	1.37	0.55
Slowed Growth		0.97	0.78	0.05
Decreased Permeability		0	0.16	0

**Notes:**

% - percent

ml – milliliter

PLFA – Phospholipid Fatty Acid Analysis

qPCR – Quantitative Polymerase Chain Reaction Analysis

SRB – sulfate reducing bacteria

<sup>1</sup>Insufficient sample volume was collected on July 16, 2014 for the full baseline sampling round, so additional sample volume was collected and submitted on July 17, 2014.**Table 2-4 Microbial Data for ST012-W30**

Sample Location:		ST012 – W30		
Parameter	Sample Type:	Baseline 7/16/2014 7/17/2014 <sup>1</sup>	Post-Shut-In 9/2/2014	Post-Extraction 9/10/2014
	Units			
qPCR				
Denitrifying Bacteria (nirK)	cells/ml	8.06E+05	4.89E+05	3.87E+05
Denitrifying Bacteria (nirS)	cells/ml	4.25E+05	3.20E+04	1.16E+04
Geobacter spp. (GEO)	cells/ml	2.33E+01	1.06E+04	1.17E+05
Sulfate Reducing Bacteria (APS)	cells/ml	4.45E+04	4.37E+04	2.00E+05
Methanogens (MGN)	cells/ml	5.05E+04	6.42E+04	4.68E+04
PLFA				
Total Biomass	cells/ml	5.75E+04	1.02E+04	4.15E+04
Firmicutes	% of Total	48.44	24.57	14.07
Proteobacteria	% of Total	22.59	26.93	50.65
Anaerobic Metal Reducers	% of Total	5.4	1.54	1.55
SRB/Actinomycetes	% of Total	3.99	4.13	0.9
General	% of Total	19.6	32.08	30.95

**Table 2-4 Microbial Data for ST012-W30**

Sample Location:		ST012 – W30		
Sample Type:		Baseline 7/16/2014 7/17/2014 <sup>1</sup>	Post-Shut-In 9/2/2014	Post- Extraction 9/10/2014
Parameter	Units			
Eukaryotes	% of Total	0	0.75	1.89
Slowed Growth		0.9	0.75	0.25
Decreased Permeability		0.12	0	0.07

**Notes:**

% - percent

ml – milliliter

PLFA – Phospholipid Fatty Acid Analysis

qPCR – Quantitative Polymerase Chain Reaction Analysis

SRB – sulfate reducing bacteria

<sup>1</sup>Insufficient sample volume was collected on July 16, 2014 for the full baseline sampling round, so additional sample volume was collected and submitted on July 17, 2014.**2.3 Solution Mixing and Injection (Injection Phase)**

Groundwater extraction, solution mixing and injection took place on 18 July 2014 for ST012-W30 and on 21 July 2014 for ST012-W11. First a submersible Grundfos Redi-Flo pump was used to extract 400 gallons from each of the test wells. Extracted water was stored in a 500-gallon tank. The extracted water was then de-oxygenated by purging with nitrogen and mixed by placing submersible pumps in the tank and recirculating the water for 30 minutes. Following de-oxygenation, 12 kilograms of potassium sulfate, the terminal electron acceptor being tested, and 300 grams of sodium bromide, the conservative tracer, were added to each of the tanks and were thoroughly mixed. A sample from each batch of solution was collected and submitted to Test America for laboratory analysis of bromide and sulfate. Table 2-5 provides the calculated concentrations of bromide and sulfate compared to the laboratory results.

**Table 2-5 Concentrations of Solution Used for Injection**

Location	Bromide Concentration (mg/l)		Sulfate Concentration (mg/l)	
	Calculated	Lab Result	Calculated	Lab Result
ST012-W11	150	150	4,300	2,300
ST012-W30	150	220	4,300	3,800

**Notes:**

mg/l – milligrams per liter

Once the test solution was mixed, the solution was gravity fed into the associated test well. A water level indicator was used to monitor groundwater levels during solution injection to make sure an overflow condition did not occur.

**2.4 Shut-In Period**

Following the push phase of the field test, samples were collected twice per week from each monitoring well using low-flow procedures and were sent to Test America for bromide and sulfate analysis. Additionally, readings were collected from the bromide meters to provide instantaneous data to enable decision making with regards to how long to wait prior to conducting the pull-phase

of the test. The data from the bromide probes, however, were deemed unreliable early on in the shut-in period. The bromide probes were removed from the wells, re-calibrated, and replaced inside the wells, but the data continued to be unreliable. On 16 August 2014, the bromide probes were removed from the wells and sent back to the supplier. Given that the bromide probes could not be used, the bi-weekly sample results were compiled, and trend graphs were created and updated as soon as data became available so that decisions could be made on when to conduct the pull phase.

As indicated in the EBR Field Test Plan, it was anticipated that the shut-in period would last between two and four weeks. The actual duration was dependent on how long it would take for the conservative tracer, bromide, to reach ambient conditions. The shut-in period would last no more than twice the duration that it took to reach ambient conditions. Based on laboratory data through 19 August 2014, approximately four weeks into the shut-in period, with concentrations approaching baseline conditions (Table 2-6), a decision was made to schedule the extraction phase for the week of 1 September 2014.

**Table 2-6 Bromide and Sulfate Concentrations during Shut-In Period**

Sample Type	Date	ST012–W11		ST012–W30	
		Bromide (mg/l)	Sulfate (mg/l)	Bromide (mg/l)	Sulfate (mg/l)
Baseline	7/16/2012	1.6	5.4	1.3	11
Shut-in	7/22/2014	55	1,000	99	1,900
Shut-in	7/24/2014	39	2,000	92	1,600
Shut-in	7/29/2014	18	940	47	840
Shut-in	7/31/2014	12	610	35	660
Shut-in	8/5/2014	6.2	300	20	320
Shut-in	8/7/2014	4.9	230	16	240
Shut-in	8/12/2014	3.0	110	11	140
Shut-in	8/15/2014	2.4	73	8.7	100
Shut-in	8/19/2014	1.9	42	6.5	67
Shut-in	8/21/2014	1.8	31	5.2	49
Shut-in	8/26/2014	1.6	18	4.5	34
Shut-in	8/29/2014	1.5	14	3.9	24
Post-Shut-In	9/2/2014	2.6	6.4	3.5 J	18

**Notes:**

mg/l – milligrams per liter

J = The analyte was positively identified; the quantitation is an estimation

Prior to conducting extraction, low flow samples were collected from each monitoring well, and were submitted to Test America and Microbial Insights to be tested for the full suite of analysis that was conducted during baseline sampling. The results of these samples are presented in Tables 2-1 through 2-4 and discussed in Section 3.

## 2.5 Extraction Phase

For the extraction phase of the field test, a submersible Grundfos Redi-Flo pump was installed sequentially in ST012-W11 and ST012-W30 at 248 and 216 feet below top of casing, respectively.

The pump extracted groundwater at a rate of up to 20 gallons per minute (gpm) to extract as much of the solution that was introduced to the groundwater as possible.

Extraction started at ST012-W11 on 4 September 2014. 1,000 gallons of water were extracted from ST012-W11 on 4 September 2014 and 9,000 gallons were extracted during a continuous period from 6 September through 7 September 2014. A pumping rate of 5 to 6 gpm was maintained during extraction. Extracted water was pumped into a 21,000-gallon frac tank and a slip line from the main conveyance line was used to collect a sample at intervals of approximately 500 gallons. The collected samples were shipped to Test America for bromide and sulfate analysis. Bromide and Sulfate results during the pull phase in ST012-W11 are tabulated in Table 2-7.

**Table 2-7 Bromide and Sulfate Concentrations for ST012-W11 during Extraction Phase**

Date / Time	Quantity Pumped (gallons)	Bromide (mg/l)	Sulfate (mg/l)	Date / Time	Quantity Pumped (gallons)	Bromide (mg/l)	Sulfate (mg/l)
9/4/14 15:05	500	2.9	260	9/7/14 9:57	5,500	2.60	280
9/4/14 16:41	1,000	2.8	260	9/7/14 11:29	6,000	2.50	280
9/6/14 21:47	1,674	2.6	320	9/7/2014 13:43	6,500	2.50	280
9/6/14 22:48	2,004	2.6	260	9/7/2014 15:25	7,000	2.50	280
9/7/14 0:23	2,500	2.6	270	9/7/2014 17:00	7,500	2.50	280
9/7/14 1:55	3,000	2.6	270	9/7/2014 18:33	8,000	2.80	290
9/7/14 3:30	3,500	2.6	270	9/7/2014 20:18	8,500	2.70	280
9/7/14 5:03	4,000	2.6	280	9/7/2014 21:54	9,000	2.70	280
9/7/14 6:35	4,500	2.6	280	9/7/2014 23:33	9,500	2.70	280
9/7/14 8:26	5,000	2.6	280	9/8/14 01:10	10,000	2.70	280

**Notes:**

mg/l – milligrams per liter.

Initial results from Test America for the pull-phase of ST012-W11 were used to calculate the total amount of sulfate that was extracted from the groundwater. The results of this calculation indicated that more sulfate was extracted from the groundwater than was introduced during the push-phase of the field test. A request was made that the laboratory re-analyze a select number of samples, and the reported results were confirmed. During re-analysis, Amec Foster Wheeler determined that the high sulfate concentrations reported by the laboratory were within range of background concentrations at the site.

On 7 September 2014, groundwater extraction began at ST012-W30 using the same techniques as at ST012-W11. Extraction of the first 500 gallons removed from this well occurred over a 14-hour period. Pumping rates from this well could not be sustained at a constant rate and required personnel to toggle the pump on and off. Due to the low yield of the well, the target volume of water to be extracted during the pull-phase was reduced to 1,000 gallons. During extraction of the remaining 500 gallons of water to be pumped from the well, three rounds of samples were collected. Extraction of this volume of water occurred over an approximate 25-hour period. The collected samples were shipped to Test America for bromide and sulfate analysis. Bromide and Sulfate results for samples collected during the pull phase in ST012-W30 are tabulated in Table 2-8. Historical groundwater monitoring upgradient of site contamination has shown background sulfate concentrations generally range from 250 to 300 milligrams per liter (mg/l) (BEM,1998) Sulfate concentrations in ST012-W30 below typical background concentrations indicate potential LNAPL-influenced conditions.

**Table 2-8 Bromide and Sulfate Concentrations for ST012-W30 during Extraction Phase**

Date / Time	Quantity Pumped (gallons)	Bromide (mg/l)	Sulfate (mg/l)
9/7/14 6:05	500	2.7	18
9/9/14 8:03	500	2.3	15
9/9/14 10:49	667	2.6	15
9/9/14 13:45	834	2.6	19
9/10/14 10:56	1,000	2.4	16

**Notes:**

mg/l – milligrams per liter

## 2.6 Post-Extraction Sampling

At the conclusion of the extraction phase for each well, post-extraction samples were collected and submitted to Test America and Microbial Insights to be tested for the full suite of analysis that was conducted during baseline sampling. The results of these samples are presented in Tables 2-1 through 2-4 and discussed in Section 3.

## 2.7 Waste Management

The groundwater purged for low flow sampling during the shut-in period and the water purged from the test wells during the pull phase of the EBR field tests were contained in a single 21,000 gallon frac tank. The groundwater was treated in the SEE groundwater treatment system.

## 2.8 Deviations from the Work Plan

Based on unforeseen conditions, the following deviations from the work plan were implemented:



- 360 1. The in-line bio-flo™ filters were not used to collect samples for microbial analysis because  
361 they clogged before the required amount of water was collected. Microbial Insights  
362 indicated that this situation occurs frequently and that collecting bottles of water that the  
363 lab will process through a filter prior to testing provides the same level of confidence with  
364 the data. Therefore, this is not a significant deviation.
- 365 2. The in-situ bromide probes did not provide reliable data and were removed from the wells  
366 within the early stages of the shut-in period. Amec Foster Wheeler relied upon laboratory  
367 data to make field decisions during the shut-in period, which required a few additional days  
368 for results to be available; trend predictions were used to schedule a time to conduct the  
369 extraction phase. Based on the results of laboratory testing, the extraction phase was  
370 conducted within the appropriate timeframe; therefore, the suspension of the use of the  
371 in-situ bromide probes is not considered a significant deviation from the work plan.
- 372 3. Due to the slow extraction rates achievable from ST012-W30, only 1,000 gallons of water  
373 was removed during the extraction phase compared to the 10,000 gallons targeted in the  
374 EBR Field Test Plan. This may be due to fouling of the well over time. Well fouling limits  
375 evaluation of hydraulic conductivity for the well. Extraction of a smaller volume of water  
376 than planned results in only partial extraction of the injected fluids. This limits evaluation  
377 of degradation kinetics; however, data from the shut-in phase is available for calculation  
378 of kinetic parameters.

### 3.0 EVALUATION OF RESULTS AND RECOMMENDATIONS

Evaluation of the analytical data obtained during the EBR Field Test and estimation of hydraulic and biological degradation parameters are presented in this section.

#### 3.1 Evaluation of Laboratory Analytical Results

Analytical concentration data for ST012-W11 presented in Table 2-1 show no significant change between the baseline and the post-shut-in period for most of the analytes evaluated. However, there is a decrease in total TPH and total VOC concentrations observed between these monitoring periods and the post-extraction sampling round. Additionally, sulfate, calcium and chloride concentrations for the post-shut-in period increased as well. These conditions were not expected and are interpreted to be a result of cleaner/background groundwater within part of the screened interval being drawn into the well rather than pulling only injected water back into the well. Historical groundwater monitoring upgradient of site contamination has shown background sulfate concentrations generally range from 250 to 300 mg/l (BEM,1998) which is similar to the concentrations observed in ST012-W11 during the pull phase.

Results for ST012-W30 presented in Table 2-2 indicate an increase in concentration for total TPH and total VOCs in both the post-shut-in sample and post-extraction sample in comparison with the baseline sample results.

#### 3.2 Evaluation of Microbial Analytical Results

Microbial results for samples collected from ST012-W11 (Table 2-3) show an increase in the quantity of sulfate reducing bacteria between the baseline and post-shut-in sampling events. Another increase in these bacteria was observed during the post-extraction sampling event, for an overall increase in density of two orders of magnitude. Geobacter (which is implicated in biological metals reduction) increased by three orders of magnitude between baseline and post-shut-in sampling rounds and then decreased by two orders of magnitude during the post-extraction sampling event. Methanogenic bacterial densities decreased by two orders of magnitude between the baseline and post-extraction sampling events.

Although increases in total biomass were modest over the course of the test, the percentages of Firmicutes and Proteobacteria were approximately equal during baseline sampling and shifted such that Proteobacteria was more prevalent in the post-shut-in sample and even more prevalent in the post-extraction sample. An increase in the proportion of Proteobacteria present coupled with an increase in biomass suggests metabolism of a readily available growth substrate. The decline in Firmicutes suggests a transition to a more aerobic environment over the course of the test and/or the extraction of more oxygenated water from upgradient of contamination (see Section 3.1).

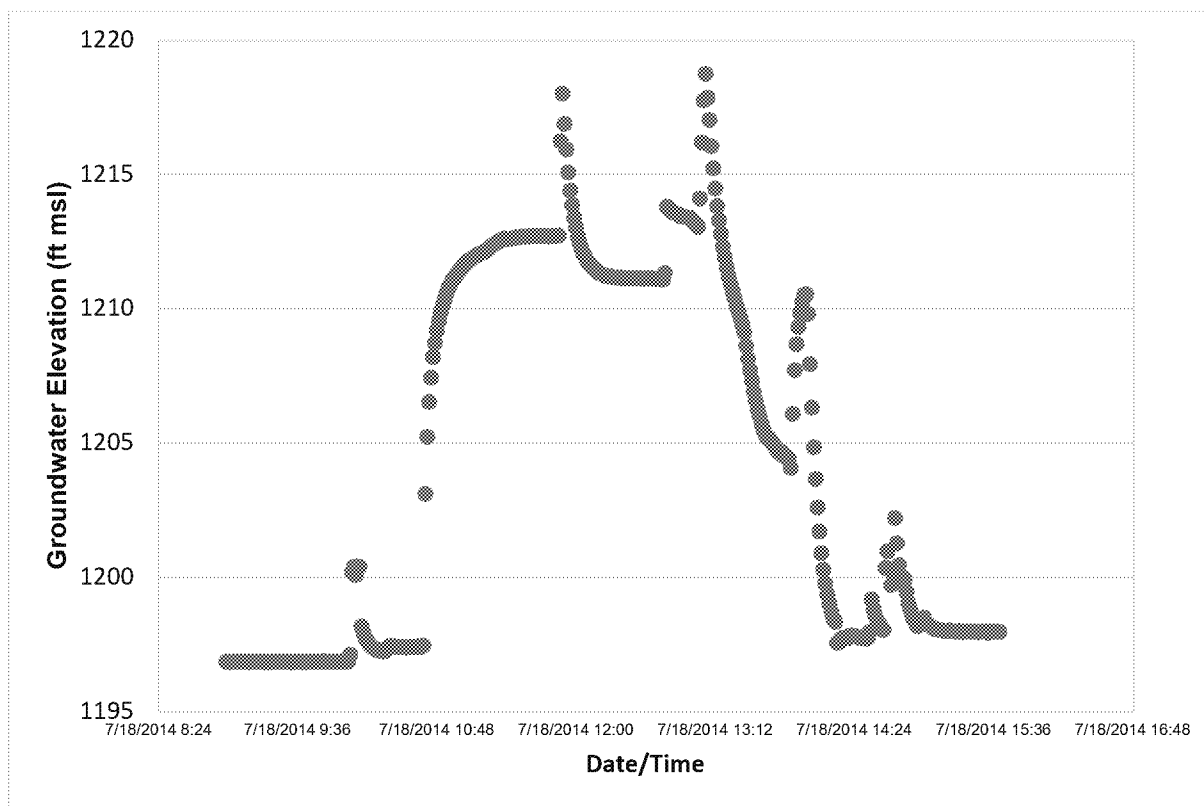
Microbial results for samples collected from ST012-W30 (Table 2-4) show no change in sulfate reducing bacteria between the baseline and post-shut-in sampling events; however, the density of these organisms increased by an order of magnitude between the post-shut-in and the post-extraction sampling events. Geobacter densities increased by three orders of magnitude between baseline and post-shut-in sampling rounds and increased by another order of magnitude during the post-extraction sampling event. Methanogenic bacterial densities decreased by two orders of magnitude between the baseline and post-extraction sampling events.

The percentage of Firmicutes was higher than the percentage of Proteobacteria in the baseline sample but shifted such that percentages of these groups were relatively the same in the post-shut-in sample; Proteobacteria were more prevalent in the post-extraction sample.

### 3.3 Estimation of Hydraulic Parameters

Water elevations from transducer data collected throughout the field test were evaluated for estimation of hydraulic parameters. However, groundwater elevation data from the transducers generally showed rapid and abrupt changes during the pull phases which was likely related to fouling of the well screens; this limited analysis of pull phase data for estimation of hydraulic conductivity. Transducer data was not recorded for the ST012-W11 push phase so only groundwater elevation data collected during the push phase in ST012-W30 provides potentially useable data. The push phase in ST012-W30 was analyzed using the groundwater model from the RD/RAWP. The purpose was to calibrate the model to the observed elevation increases associated with the ST012-W30 push phase by varying the hydraulic conductivity. The push-phase liquid injection was modeled and the predicted groundwater elevations were compared to the observed elevations from the transducer. Unfortunately, the groundwater elevation data corresponding to the push phase included about four elevation spikes related to increases in solution addition rates (see Graph 3-1) and the model could not be effectively calibrated to this data. Therefore, no refinement to the model hydraulic conductivity values are recommended based on EBR Field Test results.

**Graph 3-1 ST012-W30 Injection-Phase Elevations**



**Notes:**

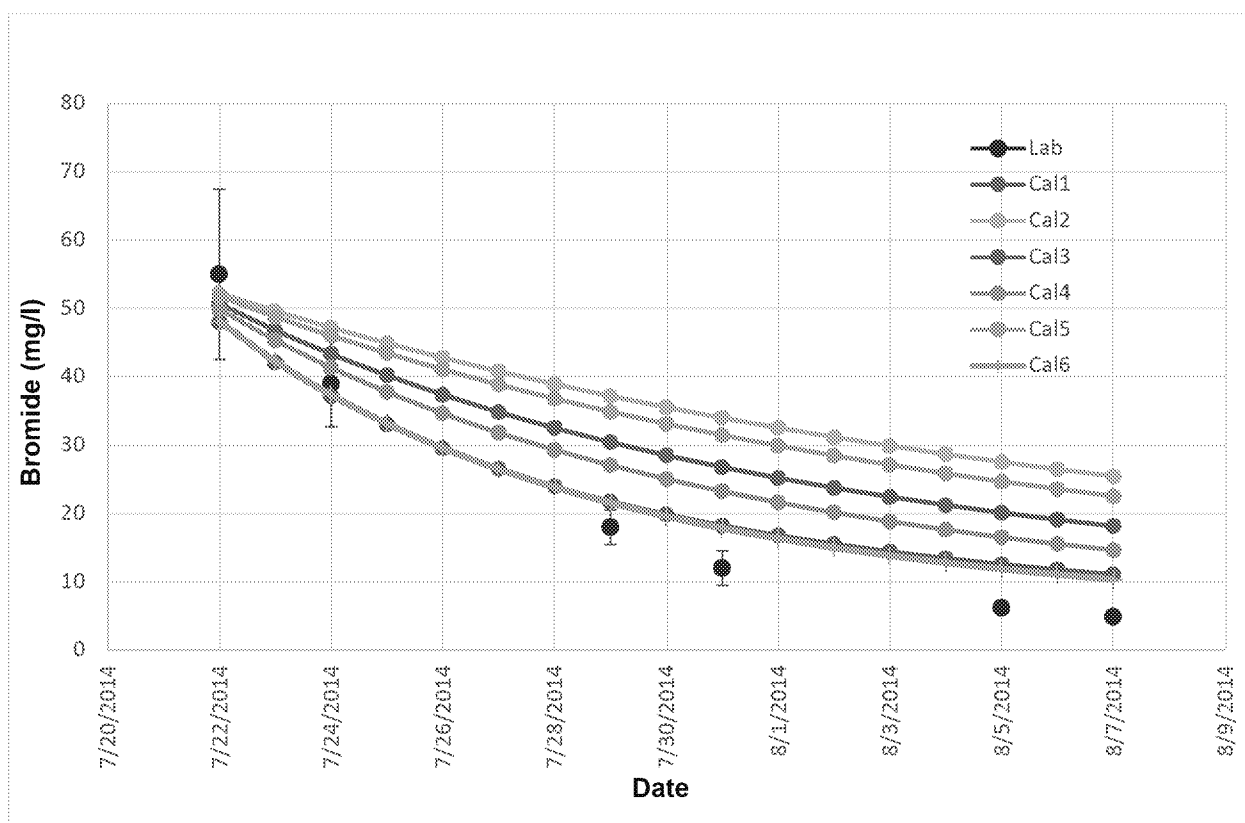
ft msl – feet above mean sea level

A similar approach was used to evaluate dispersivity at ST012 by varying dispersivity in the groundwater model to generate predicted bromide concentrations over time that compare well to the laboratory analytical results for bromide. The groundwater model uses a single set of longitudinal, horizontal, and vertical dispersivities throughout the model domain so the objective was to obtain the best calibration to both the ST012-W11 and ST012-W30 data with the same dispersivity values. Table 3-1 shows the dispersivity values for the six calibration runs and Graphs 3-2 and 3-3 show the calibration model run predicted bromide results compared to the analytical data. The shape of the bromide curve for ST012-W30 is closer to what is predicted by typical ratios of longitudinal, horizontal, and vertical dispersivities so the calibration runs focused on matching that data.

**Table 3-1 Dispersivity Values Used in Groundwater Model Calibrations**

Calibration Run	Longitudinal	Horizontal	Vertical
1	19	1.9	0.19
2	9.5	0.95	0.095
3	38	3.8	0.38
4	20	6.66	1
5	10	3.33	0.5
6	30	10	3

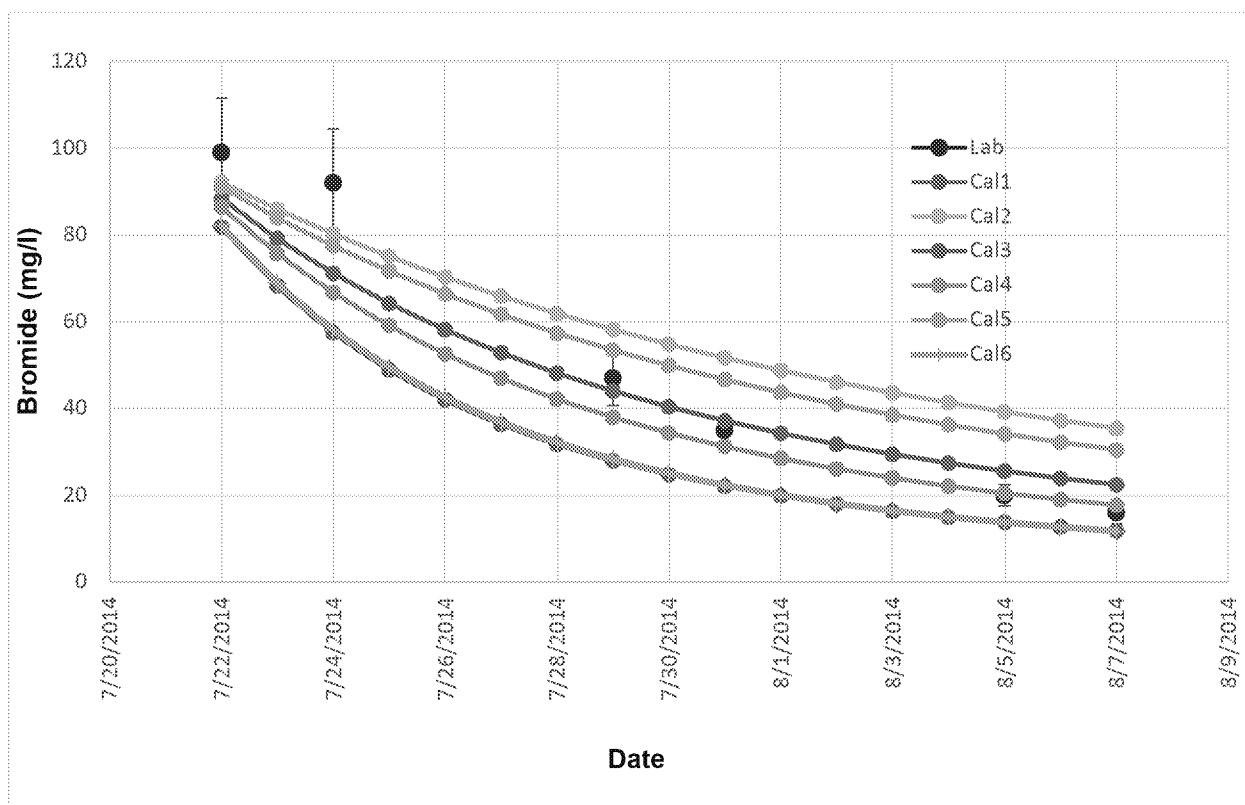
**Graph 3-2 ST012-W11 Dispersivity Calibration Runs**



**Notes:**

mg/l – milligrams per liter

461

**Graph 3-3 ST012-W30 Dispersivity Calibration Runs**

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**Notes:**

463 mg/l – milligrams per liter

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Based on this analysis, the recommended dispersivity values for future EBR modeling are from Calibration Run 4 (20, 6.66, and 1). These compare to values of 10, 1, and 0.1 used in previous RD/RAWP modeling for the LSZ. As can be seen by comparing the curves for Calibration Run 4 to Calibration Run 2 (which uses dispersivity values close to the RD/RAWP model), use of Calibration Run 4 values will predict lower contaminant concentrations with time than previous modeling.

**3.4 Estimation of Sulfate Utilization Rate**

Sulfate utilization has been estimated from the shut-in period of ST012-W11 and ST012-W30. Well fouling in ST012-W30 limited the extraction volume during the extraction-phase of the test. The limited volume extracted did not provide a sufficient data set to estimate sulfate utilization during the pull phase. As discussed in Section 3.1, the increasing sulfate concentrations in ST012-W11 during the pull phase of the test indicate that background groundwater sulfate concentrations were being pulled into the well and prevent the accurate estimation of sulfate utilization during the pull phase.

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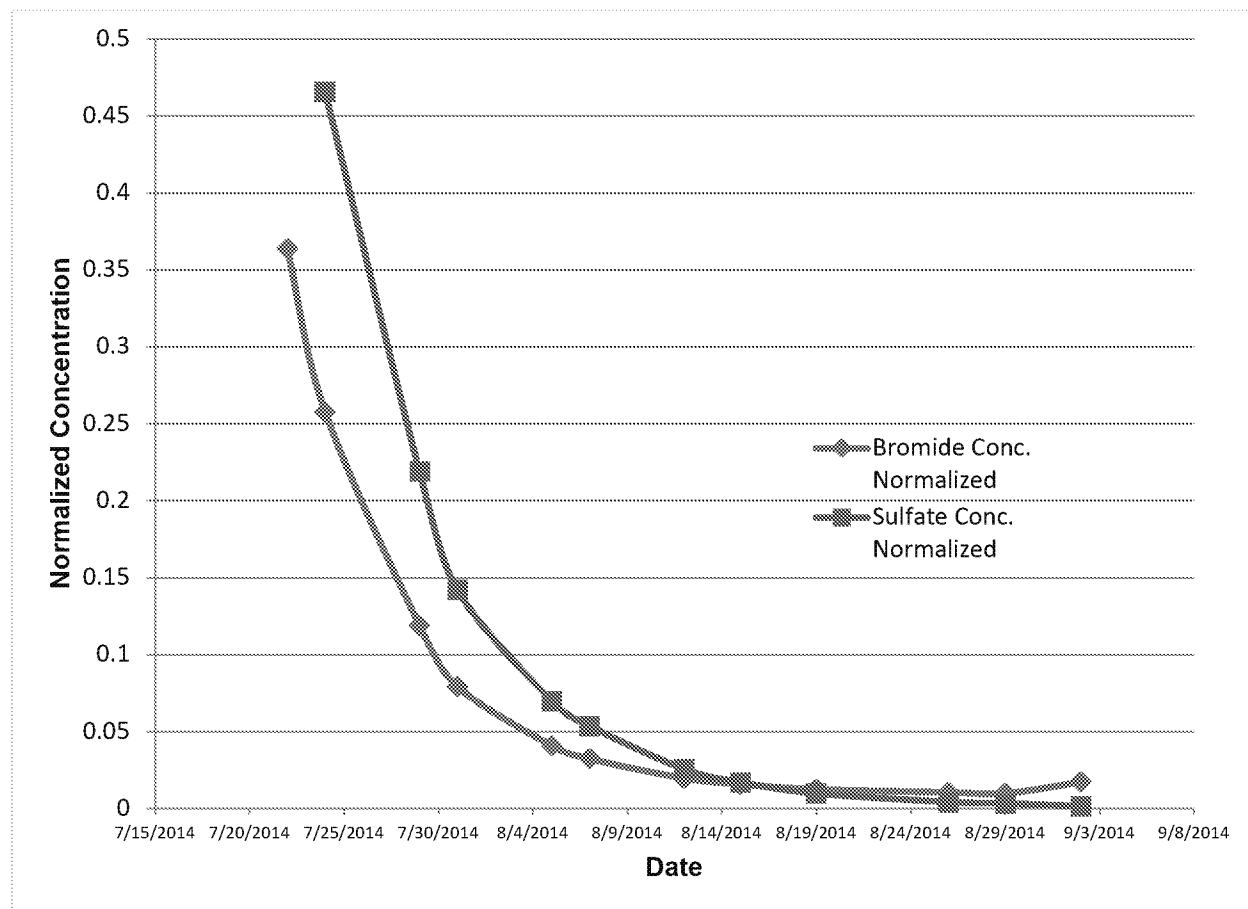
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485

Instead, sulfate utilization was estimated from sulfate concentration collected during the shut-in period. Graph 3-4 shows normalized concentrations of bromide and sulfate (measured concentration divided by initial calculated concentration) for ST012-W11 during the shut-in period. The normalized sulfate concentration is higher than the normalized bromide concentration for the majority of the shut-in period; however, after the initial 24 July 2014 sample, sulfate decreased

486 faster than bromide and the data after this date are useful for evaluating the sulfate utilization rate.  
 487

488 **Graph 3-4 ST012-W11 Shut-In Phase Bromide and Sulfate**

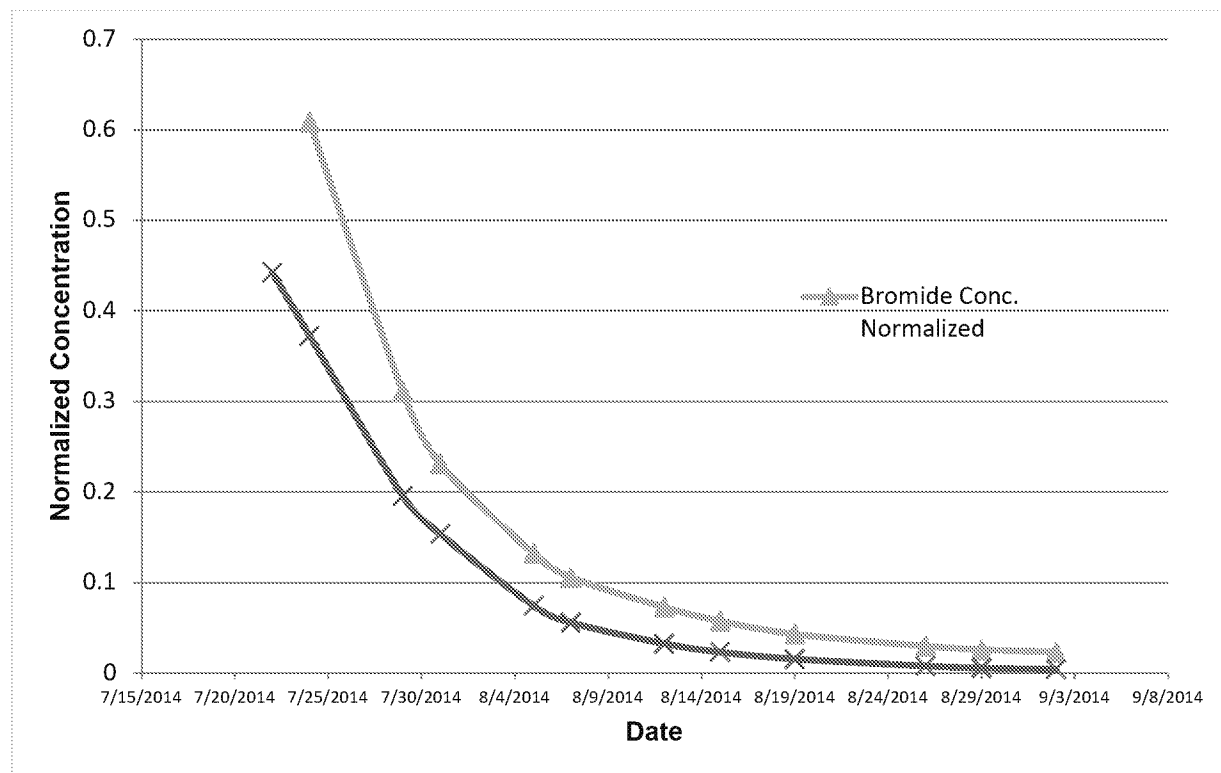


489 The data collected during the shut-in period for ST012-W30 also provides a useful data set for  
 490 estimation of the sulfate utilization rate. Graph 3-5 shows the normalized concentrations for  
 491 ST012-W30. Normalized sulfate concentrations are lower than normalized bromide  
 492 concentrations throughout the shut-in period which is expected where sulfate is being consumed.  
 493 Using the data from the ST012-W11 and ST012-W30 shut-in periods, a Lineweaver-Burk analysis  
 494 was performed. In this analysis, the half saturation coefficient and the maximum substrate  
 495 utilization rate were estimated from a graph of the inverse utilization rate and the inverse sulfate  
 496 concentration. Using this method (see calculations in Appendix E), the maximum sulfate utilization  
 497 rate ranges from 33 to 75 mg/L-day for ST012-W11 and ST012-W30, respectively. The half  
 498 saturation coefficient, which represents the sulfate concentration at which reaction rate is  $\frac{1}{2}$  of  
 499 the maximum rate, ranges from 4,100 to 4,200 mg/l of sulfate for ST012-W30 and ST012-W11  
 500 respectively.  
 501  
 502

503 These kinetic values were then used to model the sulfate consumption and TPH degradation  
 504 using Monod-type kinetics. Initial sulfate and TPH concentrations used in the model were selected  
 505 to represent background influx and LNAPL influenced conditions, respectively. A first-order  
 506 degradation curve was fitted to these modeled values to graphically approximate a first-order  
 507 maximum TPH degradation rate with a coefficient of  $0.03 \text{ day}^{-1}$  (see Appendix E). This value  
 508

compares to a value of  $0.0125 \text{ day}^{-1}$  for the maximum utilization rate of hydrocarbons other than benzene in the RD/RAWP modeling indicating that the maximum degradation rate under sulfate reducing conditions in the EBR field test was approximately 2.4 times greater than previously modeled. First order kinetics are a reasonable representation of Monod degradation kinetics when the sulfate concentration is much less than the half saturation coefficient. This is the case for background sulfate concentrations; therefore, the RD/RAWP model is based on reasonable but conservative kinetic values. Future updates to the model should incorporate the estimated Monod-type kinetic values to evaluate TPH degradation rates for sulfate concentrations approaching and greater than the half saturation coefficient.

**Graph 3-5 ST012-W30 Shut-In Phase Bromide and Sulfate**



### 3.5 Recommendations

The data collected for sulfate degradation from the EBR Field Test indicates that sulfate degrading bacteria populations increased and that dispersivity values and sulfate utilization rates were more favorable than the assumed values used in the RD/RAWP EBR modeling. These findings, in combination with previous studies that concluded sulfate reduction was the dominant naturally occurring process for contaminant assimilation (BEM, 1998), indicate that sulfate amendment should be included in the EBR strategy. Sulfate amendment could either be used solely or in combination with aerobic methods to achieve remediation goals.

The next step will be to develop a refined EBR approach using the new parameters developed in this study and to prepare an RD/RAWP addendum to document the revised approach and implementation strategy.

## 4.0 REFERENCES

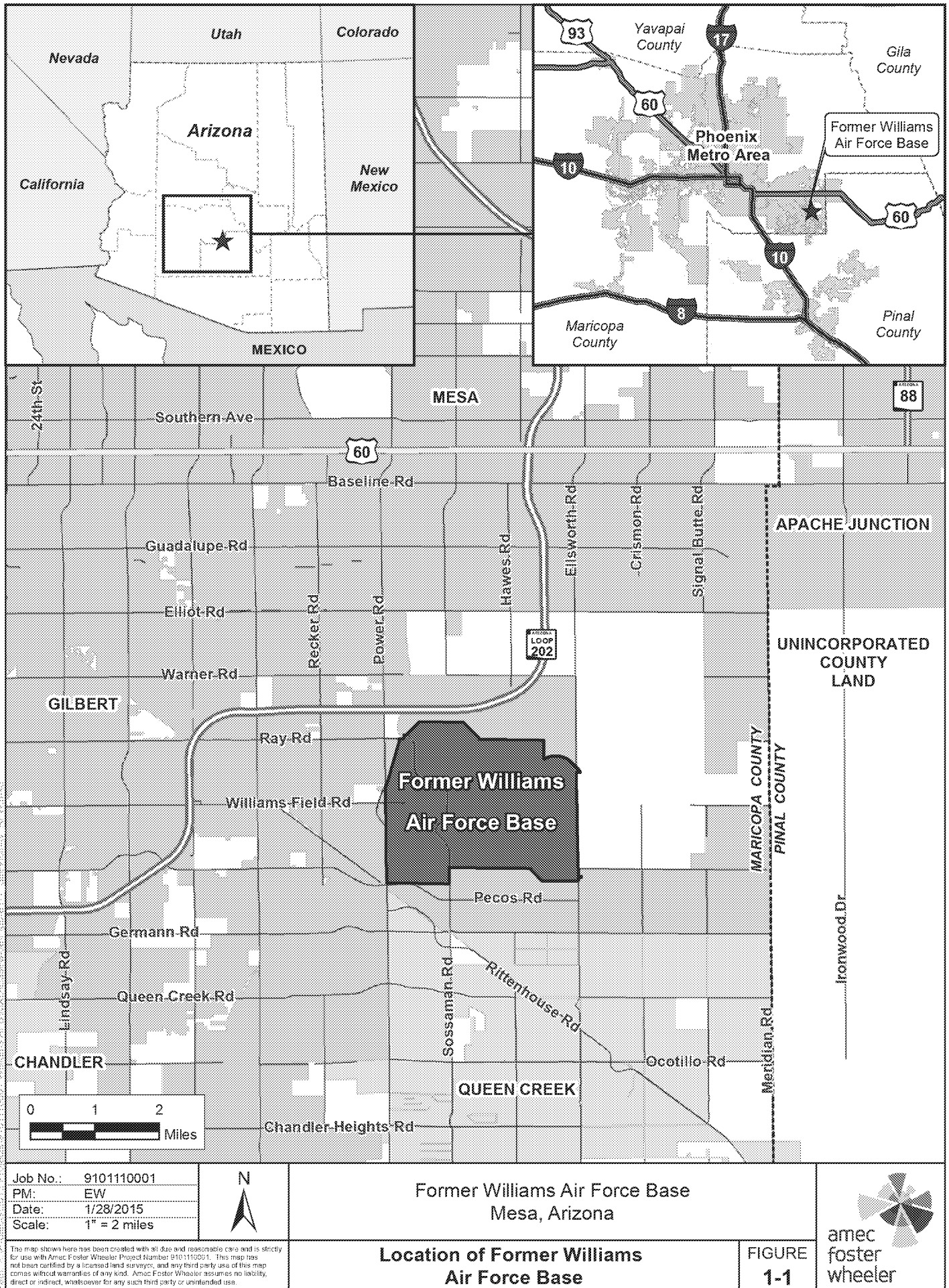
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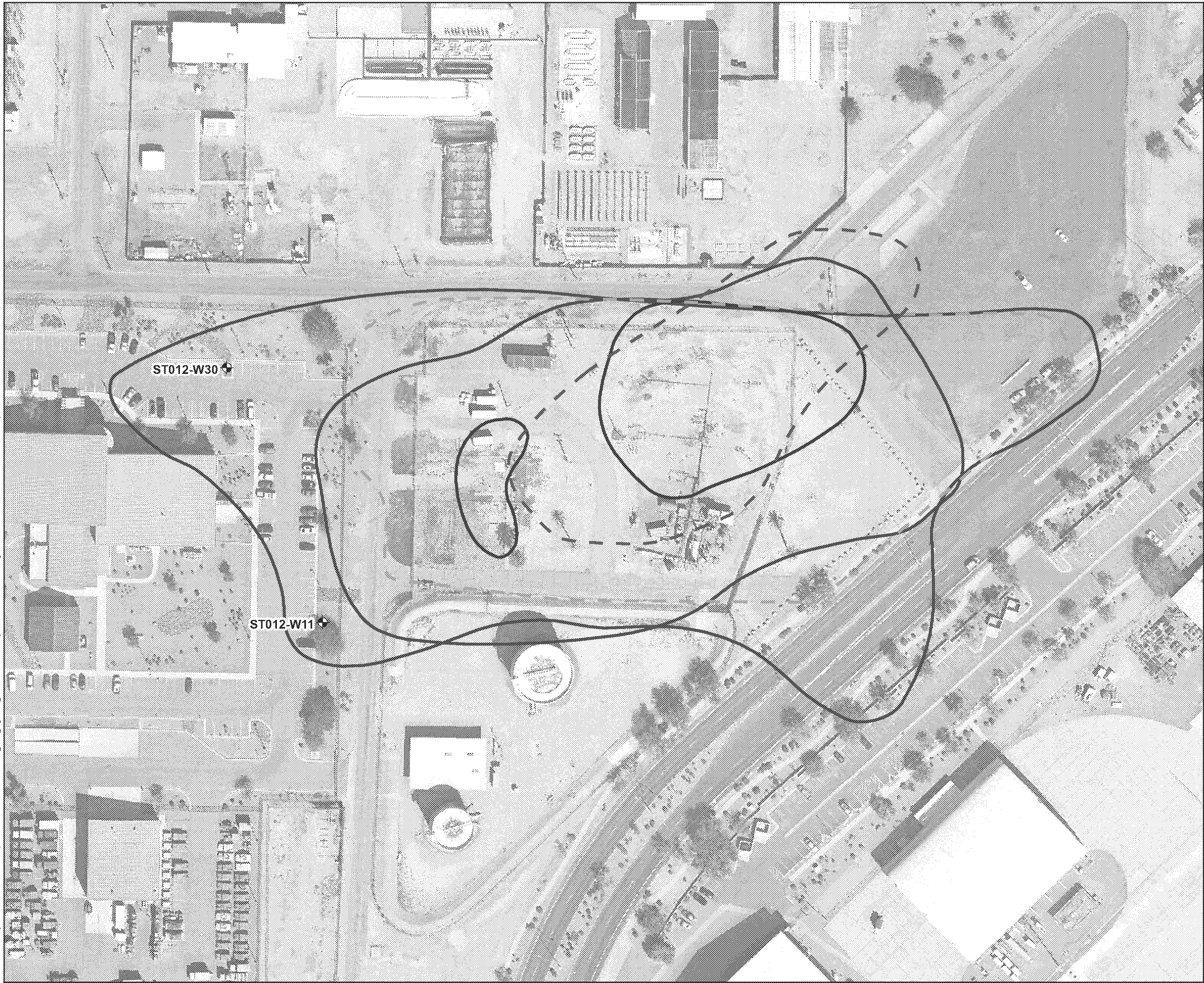
## FIGURES

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**Legend**

- Push-Pull Test Well
- CZ RD/RAWP LNAPL extent
- UWBZ RD/RAWP LNAPL extent
- LSZ RD/RAWP LNAPL extent
- LSZ SEE Treatment Zone
- CZ/UWBZ SEE Treatment Zone

Notes: SEE - Steam Enhanced Extraction  
CZ - Cobble Zone  
LSZ - Lower Saturated Zone  
UWBZ - Upper Water Bearing Zone  
LNAPL - Light Non-Aqueous Phase Liquid  
EBR - Enhanced Bioremediation  
RD/RAWP - Remedial Design/Remedial Action Work Plan



Draft Enhanced Bioremediation Field Test Report Former Williams Air Force Base Mesa, Arizona	
EBR FIELD TEST LOCATIONS	FIGURE 1-3
Job No. PM: 9101110001 Date: 4/22/2015 Scale: 1" = 100 feet	 amec foster wheeler
<small>The map shown here has been created with all due and reasonable care and is strictly for use with AMEC Project Number 9101110001. This map has not been certified by a licensed land surveyor, and any third party use of this map comes without warranties of any kind. AMEC assumes no liability, direct or indirect, whatsoever for any such third party or unintended use.</small>	

## **APPENDIX A**

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### **SAMPLE COLLECTION RECORDS**